



Docket No.: 56446-20005.20/
007006 / D1150-6US
(PATENT)

IN THE UNITED STATES PATENT AND TRADEMARK OFFICE

In re Patent Application of:
David LAM et al.

Application No.: 09/888,224

Confirmation No.: 8097

Filed: June 22, 2001

Art Unit: 1634

For: ENDOGLUCANASES, NUCLEIC ACIDS
ENCODING THEM AND METHODS FOR
MAKING AND USING THEM (AMENDED)

Examiner: Jehanne Sitton

MS Amendment
Commissioner for Patents
P.O. Box 1450
Alexandria, VA 22313-1450

DECLARATION UNDER 37 C.F.R. § 1.132

Sir:

1. I, Jay Short, am a co-inventor with David E. Lam and Eric J. Mathur, on the above-identified patent application.
2. I am an expert in the field of molecular biology and enzyme development and was an expert at the time of the invention. I am presently employed as CEO and as a research scientist at Diversa Corporation, San Diego, CA, assignee of the above-referenced patent application. My resume is attached as documentation of my credentials.
3. I declare that at the time of the invention it would not have required any knowledge or guidance as to which specific structural elements (including, e.g., domain structures, location active sites, interaction with co-factors or regulatory molecules, secondary and tertiary

structure) correlate with endoglucanase activity to create variants of the exemplary endoglucanase of the invention and test the variants for a desired endoglucanase activity. It would not have been necessary for the skilled artisan to know or predict beforehand which specific regions or structural elements of an endoglucanase were necessary for function or activity to routinely generate the genus of endoglucanase-encoding nucleic acids of the invention. At the time of the invention, methods for making and screening endoglucanases were sufficiently comprehensive and routine to predictably generate a genus of endoglucanase-encoding sequences without need of knowing which specific regions or structural elements of a sequence or structure affected function or activity. Methods known at the time of the invention for modifying nucleic acid and polypeptide sequences in combination with high through-put enzyme (e.g., endoglucanase) screening assays made methods that required previous knowledge of structural elements necessary for enzymatic activity obsolete and unnecessary. High through-put enzyme screening methodologies known at the time of the invention (including *in vivo* and *in vitro* nucleic acid expression and enzyme (endoglucanase) screening protocols) made methods that required previous knowledge of protein structure, including secondary or tertiary structure, active site sequences, and the like obsolete and unnecessary. Using methods known in the art at the time of the invention it would not have been necessary to understand which specific regions of enzyme structure (e.g., domain structures, active sites, secondary and tertiary structure) could, or could not, be modified to generate the genus of nucleic acids of the invention without undue experimentation. The specification provided sufficient guidance to one of ordinary skill in the art to make and use the genus of endoglucanase-encoding nucleic acids to practice the invention.

4. However, if the skilled in the art at the time of the invention elected to use elements of enzyme structure for guidance in designing and making variants, using the teaching of the specification the artisan had many sources of direction to understand the structure of endoglucanases to have direction and guidance in determining which amino acid residues could be substituted, deleted or inserted into a nucleic acid to obtain structural and functional variants of an endoglucanase. For example, the specification provides guidance as to what base and residue changes could be made to make the genus of endoglucanase-encoding nucleic acids of the

invention; see, for example, the paragraph from line 31, page 10 to line 16, page 11, and, page 51, lines 16 to 24, of the specification. Direction to the skilled artisan as to which amino acid residues can be substituted, deleted or inserted into a nucleic acid to obtain structural, and functional homologues of an enzyme could also be found in the art at the time of the invention. For example, Dominguez (1996) "The crystal structure of a family 5 endoglucanase mutant in complexed and uncomplexed forms reveals an induced fit activation mechanism," J. Mol. Biol. 257(5):1042-1051, describes the crystal structure of an endoglucanase; Ducros (1995) "Crystal structure of the catalytic domain of a bacterial cellulase belonging to family 5", Structure 3(9):939-49, describes the crystal structure of the catalytic domain of an endoglucanase; Davies (1995) "Structures of oligosaccharide-bound forms of the endoglucanase V from *Humicola insolens* at 1.9 Å resolution," Biochemistry 34(49):16210-20, describes the crystal structures of an endoglucanase in various forms; to name only a few examples.

I hereby declare that all statements made herein of my own knowledge are true and that all statements made on information and belief are believed to be true; and further that these statements were made with the knowledge that willful false statements and the like so made are punishable by fine or imprisonment, or both, under Section 1001 of Title 18 of the United States Code and that such willful false statements may jeopardize the validity of the application or any patent issued thereon.

Respectfully submitted

Date: _____

Jay Short

CURRICULUM VITAE

NAME Jay M. Short, Ph.D.

Dr. Short is a founding member of Diversa Corporation, has served as Chief Technology Officer and Director of the company since its inception in 1994. He assumed the additional roles of President in 1998 and Chief Executive Officer in 1999. In February of 2000, Dr. Short led the company's highly successful initial public offering, which raised over \$200 million in gross proceeds – the largest biotechnology IPO ever completed at the time. Diversa was recently named one of the 100 most influential companies that will have the greatest influence on the future of human health. Diversa Corporation (NASDAQ: DVSA) is a leader in applying proprietary genomic technologies for the rapid discovery and optimization of novel products from genes and gene pathways.

EDUCATION

| | |
|-------------|--|
| 2003 | Certified Director Director Training Program The Anderson Graduate School of Management, University of California, Los Angeles |
| 1981 - 1985 | Ph.D., Biochemistry Case Western Reserve University, Cleveland, Ohio |
| 1980 - 1981 | Graduate Study, Macromolecular Science Case Western Reserve University, Cleveland, Ohio |
| 1976 - 1980 | B.A. with Honors, Chemistry Taylor University, Upland, Indiana |

RESEARCH & PROFESSIONAL EXPERIENCE

| | |
|----------------|--|
| 1999 - present | CEO and President Chief Technology Officer Board of Director Diversa Corporation San Diego, California |
| 1998 - present | President and Chief Technology Officer Board of Director Diversa Corporation San Diego, California |
| 1997 - 1998 | Executive Vice President and Chief Technology Officer Board of Director Diversa Corporation San Diego, California |
| 1994 - 1997 | Chief Technology Officer Board of Director Diversa Corporation San Diego, California |
| 1990 - 1994 | President Stratacyte, Inc. La Jolla, California |

Jay M. Short, Ph.D.

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|-------------|---|
| 1992 - 1994 | Vice President R&D (Research) and Operations Stratagene Cloning Systems La Jolla, California |
| 1989 - 1992 | Vice President R&D (Research) and Biological Operations Stratagene Cloning Systems La Jolla, California |
| 1988 - 1989 | Senior Staff Scientist Research and Development Stratagene Cloning Systems La Jolla, California |
| 1985 - 1988 | Staff Scientist Research and Development Stratagene Cloning Systems La Jolla, California |
| 1981 - 1985 | Ph.D. Research Case Western Reserve University Dr. Richard W. Hanson's Laboratory, Identification and characterization of the promoter for P-enolpyruvate carboxykinase. First identification of a cAMP regulatory domain. Cleveland, Ohio |
| 1980 - 1981 | Graduate Student Research Case Western Reserve University Dr. Bruce Roe's Laboratory, Analysis of the cellulase activity of <i>Trichoderma viride</i> . Cleveland, Ohio |

TEACHING EXPERIENCE

Thesis Advisor, University of Uppsala, Sweden, Ph.D. for Michelle Alting-Mees 1988-1993
Lecturer, Committee for Advanced Scientific Education, Center for Drug Evaluation and Research, FDA 1992
Faculty, Transgenic Mouse Model and Its Application in Assessing *In Vivo* Mutagenesis, Genetic
Toxicology Workshop (3rd Annual) 1989
Microbiological Associates Inc., Bethesda, MD.
Faculty, DNA Cloning and Expression, Physiology Society Workshop, San Diego, CA. 1987
Teaching Assistant, Molecular & Cellular Biology, Case Western Reserve University, Cleveland, OH. 1981-1985
Teaching Assistant, Physiological Chemistry, Kent State University, Kent, OH. 1981
Teaching Assistant, Quantitative Analysis, Taylor University, Upland, IN. 1978-1980

CERTIFICATIONS

| | |
|--------------------|--|
| Certified Director | Director Training Program, University of California, Los Angeles, California The Anderson Graduate School of Management and The Harold Price Center |
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for Entrepreneurial Studies

PADI Diver Certification

PROFESSIONAL EXPERIENCE

Diversa ranked # 2 among small companies for one of the best places for life scientists to work in this industry.
Diversa named one of the 100 most influential companies that will have the greatest influence on the future of human health, Acumen 2004

Diversa's patent portfolio ranked # 1 on the 2003 Patent Scorecard by the MIT Survey

Largest Biotechnology IPO raising over \$200MM

Founding management member of Diversa Corporation

Board Director, Diversa Corporation, San Diego, CA

Board Director, Invitrogen Corporation, Carlsbad, CA

Board Director, Stressgen Biotechnologies, Vancouver, Canada and San Diego, CA

Board Director, Senomyx Corporation, San Diego, CA

Board Director, YPO (Young Presidents' Organization), San Diego, CA

Board Director & Treasurer, Stressgen Therapeutics, Victoria, BC, Canada

Board Director & Secretary, Stressgen Therapeutics, Victoria, BC, Canada

Board Director & Compensation Chairman, Victoria, BC, Canada

Board Member Advisor, Chemical and Engineering News

Board Member, BioCom San Diego

Board Advisor, IngleWood Ventures

Board of Advisors and Founding Member, Division of Biological Sciences, UCSD

Board Director and Executive Committee, Zymetrics

Fellow, Lifetime, The Explorers Club, New York, NY

Committee Member BioCom Science & Technology, San Diego

Consultant, Stratagene Cloning Systems, La Jolla, CA

Consultant, Micro Product Systems, Lynn, IN

Consultant for European Economic Community on Transgenic Toxicology Testing 1991-1994

Chairman, Discussion Group, Society of Toxicology Conference 1993

Editor, Mutation Research

Judge on the U.S. National Entrepreneur of the Year 2003

Institutional Animal Care and Use Committee (IACUC), Chairman and Institutional Official

NIEHS Peer Review Committee

Panel Member for Chemical Science & Technology for NIST, National Research Council 1997-2000

SBIR Study Section

Reviewer for U.S. Congressional Office of Technology Assessment (OTA) on The Human Genome Project and Patenting DNA Sequences.

Reviewer for Proceedings of the National Academy of Sciences, Genetic Analysis Techniques, Analytical Biochemistry & Nucleic Acids Research

U.S. Committee Member for Evaluation of Biotechnology Research in Spain

Visiting Scientist, International Centre of Insect Physiology and Ecology (ICIPE), Kenya 2002-2004

MEMBERSHIPS

American Association for the Advancement of Science

American Chemical Society

American Men and Women of Science

American Society of Biochemistry and Molecular Biology

Jay M. Short, Ph.D.

American Society of Microbiology
BioCom San Diego
Environmental Mutagenesis Society
Japanese Environmental Mutagen Society
Science
Society for Industrial Microbiology
Society of Toxicology
The Explorers Club, Fellow Lifetime Member, New York
The New York Academy of Sciences
YPO (Young Presidents' Organization) San Diego
YPO (Young Presidents' Organization) International

AWARDS

Henry F. Whalen, Jr. Award for Business Development, American Chemical Society, 2004
Distinguished Alumnus Award for Professional Achievement, Taylor University, Upland, IN 2004
Taylor University nomination for CCCU Award (Council for Christian Colleges & Universities) 2003
Case Western Reserve University Alumni Profile 2003
bioFusion 03 Breakthrough Innovation in Science Award Nomination 2003
bioFusion 03 Life Science Leader of the Year Nomination 2003
bioFusion 03 Life Science Company of the Year Nomination 2003
ABL (Adaptive Business Leader) Innovations in HealthCare Gold Award 2003
Deloitte and Touche's Orange County / San Diego Technology "Fast 50" 2003
Finalists for UCSD Connect's Most Innovative New Product Award in the Biotechnology R&D Category 2002
Deloitte and Touche "Fast 500" Technology 2002
Deloitte and Touche's Orange County / San Diego Technology "Fast 50" 2002
The Premier Print Award, Annual Report 2002
Deloitte and Touche "Fast 500" Technology 2001
Ernst & Young San Diego Entrepreneur of the Year 2001
bioFusion 01 Life Science Innovator Award Nomination 2001
T-Sector Life Science Innovator Award 2001
Deloitte and Touche's Orange County / San Diego Technology "Fast 50" 2001
San Diego Business Journal StarCom Honor 2001
League of American Communication Professionals, Platinum Award, Annual Report 2001
Ernst & Young Finalist for San Diego Entrepreneur of the Year in 2000
The Premier Print Award, Annual Report 2001
American Men and Women of Science 1995
Who's Who Registry of Business Leaders 1994-1995
SBIR Annual Report Program Success Profile (Top 8 of 800 Companies) 1993
Stratagene Most Innovative Award - Managers/Supervisors 1992
Stratagene Innovation Award - Big Blue[®] Transgenic Testing System 1991
UCSD Connect Program 1st Place Award for Innovation and Entrepreneurship in Biotechnology 1991
UCSD Connect Program 1st Place Award for Innovation and Entrepreneurship in Biotechnology 1990
Stratagene Innovation Award - Lambda ZAP[®] vector 1990
Stratagene Service Award 1990
Award from the University of Victoria for Contributions to the Development of Short-term
Transgenic Mutation Assays
Nominated as Council Member for the U.S. Environmental Mutagen Society
PNIT Patent Award

MEDIA:

ABC Discovery News, ABC San Diego Channel 10, Agricultural Genomics, BBC Radio, Billings Gazette, BioCentury, Bioinformed Newsletter, BioPeople Magazine, BioTech Today Radio Show, Biotechnology Newsletter, BioVentures View, BioWorld Today, Business Daily, Business Week, CBS MarketWatch Weekend, CEO Cast, Chemical Engineering, Chemical Week, Chemistry & Industry (UK), Chemistry, CNBC, CNN Science & Technology, CNN Sunday Weekend, CNN WorldView, dBusiness.com, Digital Jam, Discovery Magazine, Drug Discovery Today, Elsevier Science Ltd., Forbes, Forbes.com, Fox CONNECT, Fox 6 News San Diego, German RTL TV, Good Morning America, Horizon Air Magazine, Idea TV, Inside Business Radio Show, JAG Financial News, KCRA Channel 3, KBPS Radio, KFMB Channel 8, KGTV Channel 10, KPBS, KUSI, Life Technology, London Financial Times, Los Angeles Times, Modern Drug Discovery, NBC San Diego Channel 7/39, National Geographic, National Radio Report, Nature, Nature Biotechnology, New York Times, PBS, Pirateinvestor.com, R&D Magazine, Reuters, San Diego Business Journal, San Diego Business Transcript, San Diego Magazine, San Diego Metropolitan, San Diego Union Tribune, SIM, Scientist, Specialty Chemicals, Sp2 Magazine, Stewards' Watch, T-Sector Magazine, The Age Magazine, The Economist, The Motley Fool, The Discovery Channel, The Discovery Channel, Time Magazine, USA Today, Wall Street Journal, Wall Street Transcript, Washington Post

PATENTS

The Patent Scorecard for 2003 recognized Diversa's patent portfolio as being ranked # 1 by the MIT Survey. This ranking provides an overall assessment of a company's intellectual property power. This measure showcases the broader significance of a company's patents by examining how often its U.S. patents from the previous five years are cited as prior art in the current year's batch. A value of 1.0 represents average citation frequency, so, for example, a value of 1.4 would indicate a company's patents were cited 40 percent more often than the average. Diversa has a value of 14.43.

DNA Cloning Vectors with *in vivo* Excisable Plasmids 1987
Mutagenesis Testing Using Transgenic Animals Carrying Marker Genes 1987
Mutagenesis Testing Using Transgenic Non-Human Animals Carrying Test DNA Sequences 1987
Dietary and Hormonal Regulation of Expression of Exogenous Genes in Transgenic Animals Under Control of the Promoter of the Gene
Phosphoenolpyruvate Carboxykinase 1988
A Transgenic Mouse for Measurement and Characterization of Mutation Induction *In Vivo* 1989
Rapid Screening Mutagenesis and Teratogenesis Assay 1989
A Combinatorial Approach to Regenerating the Immunoglobulin Repertoire in Prokaryotic Cells 1990
Transgenic Animal Models for *In Vivo* Mutagenesis Testing 1990
Polycos Vectors 1991
A Lambda Packaging Extract Lacking β -Galactosidase Activity 1991
A System for Regulation of Eukaryotic Genes 1991
Methods for Phenotype Creation from Multiple Gene Populations 1991
Transgenic Non-Human Animals Carrying Test DNA Sequences 1992
Mutagenesis Testing Using Transgenic Non-Human Animals Carrying Test DNA Sequences 1992
Selectable System Patent 1992
Polycos Mutagenesis Systems 1993
Use of Trans-acting Proteins for the Development of an *In Situ* Expression Screening System 1993
Enzyme Kits and Libraries 1995
Enzyme Activity Screening of Clones having DNA from Uncultivated Microorganisms 1995
Enzyme Tiered 1995
Method for Screening for Enzyme Activity 1995
Combined Enzyme Screening/Evolution 1995
Uncultured/Activity Screening 1995
Directed Evolution of Thermophilic Proteins 1995
Combinatorial Enzyme Development (Directed Mutagenesis) 1996
Protein Activity Screening of Clones having DNA from Uncultivated Microorganisms 1996
Production and Use of Normalized DNA Libraries 1996

Methods of DNA Shuffling with Polynucleotides Produced by Blocking or Interrupting a Synthesis or Amplification Process 1996
Method of Screening for Enzyme Activity (Biopanning) 1996
Directed Evolution of Thermophilic Enzymes 1996
Environmental Biopanning 1996
Combinatorial Enzyme Development 1996
Enzyme Activity Screening of Clones Having DNA from Uncultivated Microorganisms 1996
Normalized Samples/Libraries 1996
Reassembled Pools of Mutagenized DNA & Procedure 1996
Fluorescent-based Single Screening for Enzymes 1996
High Throughput Screening for Novel Enzymes 1997
Nucleotide Sequence of the *Aquifex aeolicus* Genome, Fragments Thereof, and Uses Thereof 1997
Screening for Novel Bioactivities 1997
Screening for Novel Compounds which Regulate Biological Interactions 1997
Method for Screening Enzyme Activity 1997
High Throughput Screening for Novel Enzymes 1997
"Discovery" (whole process, including uncultivated, normalized, biopanning, screening, evolving, (etc.) 1997
Production of Enzymes Having Desired Activities By Mutagenesis 1999
Protein Activity Screening of Clones Having DNA from Uncultivated Microorganisms 1999
Method of DNA Reassembly by Interrupting Synthesis 1999
Production and Use of Normalized DNA Libraries 1999
Enzyme Kits and Libraries 1999
Screening for Novel Bioactivities 2000
Method for Screening for Enzyme Activity 2000
Screening for Novel Bioactivities 2000
Production and Use of Normalized DNA Libraries 2000
Method of Screening for Enzyme Activity 2000
Screening Methods for Enzymes and Enzyme Kits 2001
Saturation Mutagenesis in Directed Evolution 2001
High Throughput Screening for Novel Enzymes 2001
Recombinant Bacterial Phytases and Uses Thereof 2001
Methods Useful for Nucleic Acid Sequencing Using Modified Nucleotides Comprising Phenylboronic Acid 2001
End Selection in Directed Evolution 2001
Gene Expression Library Produced From DNA From Uncultivated Microorganisms and Method for Making the Same 2001
Directed Evolution of Thermophilic Enzymes 2002
Method for Screening for Enzyme Activity 2002
Exonuclease-Mediated Gene Assembly in Directed Evolution 2002
End Selection In Directed Evolution 2002
Exonuclease-Mediated Gene Assembly in Directed Evolution 2002
Screening for Novel Bioactivities 2002
Method of DNA Shuffling with Polynucleotides Produced or Blocking or Interrupting Synthesis or Amplification Process 2002
Production and Use of Normalized DNA Libraries 2002
Sequence Based Screening 2002
Non-Stochastic Generation of Genetic Vaccines 2002
Altered Thermostability of Enzymes 2003
Screening Methods for Enzymes and Enzyme Kits 2003
Methods for Identifying a Desired Enzymatic Activity 2003
Enzymes Kits and Libraries 2003
Method for Screening for Enzyme Activity 2003
Protein Activity Screening of Clones having DNA from Uncultivated Microorganisms 2003
High Throughput Screening of Mycelia for Bioactivities of Biomolecules 2003
Screening for Novel Bioactivities 2003
Coated Surfaces for Selective Enrichment of Microbial Populations 2003

Recombinant Bacterial Phytases and Uses Thereof 2003
Synthetic Ligation Reassembly in Directed Evolution 2003
Process for Generating Optimized Molecules from a Manmade Library of Polynucleotides made by Combinatorial Saturation Mutagenesis (amended) 2003
Exonuclease-Mediated Nucleic Acid and Reassembly in Directed Evolution 2003
Methods for Purifying Annealed Doubled-Stranded Oligonucleotides Lacking Base Pair Mismatches 2004
End Selection in Directed Evolution 2004
Protein Activity Screening of Clones having DNA from Uncultivated Microorganisms 2004
Method of Screening for Enzyme Activity 2004
Exonuclease-Mediated Gene Assembly in Directed Evolution (3/23/04 new issuance) 2004
Directed Evolution of Thermophilic Enzymes (3/30/04 new issuance) 2004
Non-Stochastic Generation of Genetic Vaccines and Enzymes 2004
Directed Evolution of Thermophilic Enzymes 2004
Over 350 Additional Pending Patent Applications Worldwide.

GRANTS AND CONTRACTS

*Phase I Small Business Contract #N43-Am-62282. 1985-1986 P.I.
Vectors and Techniques for Rapid DNA Sequencing
*Phase II Small Business Contract #N43-Am-62282. 1988-1990 P.I.
Vectors and Techniques for Rapid DNA Sequencing
*Phase I Small Business Grant 2R43ES04484-02. 1986-1987 P.I.
Identification of Genetic Lesions Leading to Mutations
*Phase II Small Business Grant 2R43ES04484-02. 1989-1992 P.I.
Identification of Genetic Lesions Leading to Mutations
*1R01-ES04728-01A1. 1989-1992. (NIEHS) P.I.
Animal Model for Identification of Genetic Lesions
*Phase I Small Business Grant #R43GM42291-01. 1989 P.I.
Switch Mechanism for Gene Expression in Transgenics
*RFP NIH-ES-88-11. 1989-1994. (NIEHS) Co-I.
Development of Mutagenesis Assays Using Transgenic Mice
*Phase II Small Business Grant #2R44GM42291-02. 1990-1992 (DRG/NIH) P.I.
Switch Mechanism for Gene Expression in Transgenics
*Phase I Small Business Grant #1R43GM46585-01. 1991 (DRG/NIH) P.I.
Generation of a Peptide Screening System Through the Development of
Combinatorial-splicing "Polycos" Vectors
*Phase I Small Business Grant #1R43CA57066-01. 1992 (NCI) P.I.
Transgenic Rats: A Short-term Mutagenicity Assay for Multi-species Testing of Suspected Human Carcinogens
*Phase I Small Business Grant #1R43GM48300-01. 1992. (DRG/NIH) P.I.
Analysis of the Immunoglobulin Hypermutator Mechanism
*Phase I Small Business Grant #1R43ES06146-01. 1992 (NIEHS) P.I.
Selectable "Polycos" Shuttle Vectors for In Vivo Mutagenicity Testing
*Phase II Small Business Grant #2R44GM46585-02. 1992-1994 (NIGMS) P.I.
Peptide Screening Utilizing Combinatorial Polycos Vector
*Phase I Small Business Grant #1R43RR08667-01. 1992-1993 (DRG/NIH) Co-I.
A One-step PCR Cloning System Based on FLP Recombination
*Phase II Small Business Grant #2R44CA57066-02. 1993-1995 (NCI) P.I.
Transgenic Rats: Transgenic Rat Model for Mutagenicity Testing
*Phase I Small Business Grant. 1993-1994 (NIH) Co-I.
Transgenic Fish Model for Mutagenicity Testing
*Phase II Small Business Grant 1994-1996 (NIH) P.I.

Jay M. Short, Ph.D.

"Polycos" Shuttle Vectors for Mutagenicity testing
*Phase I Small Business Grant. 1994 (NIH) Co-I.
Vector System for Studying Protein-Protein Interactions
*CRADA with LLNL. 1994 (NIH) Co-I.
Mouse and Rat Painting Probes
*CRADA with FDA. 1994 (NIH) Co-I.
Tamoxifen Testing in F-344 Rats
*CRADA with NASA. 1994 (NIH) Co-I.
Radiation Damage in the Microgravity Environment

ABSTRACTS AND INVITED LECTURES:

Over 200 Abstracts and Invited Lectures.

PUBLICATIONS:

1. Yoo-Warren, H., Monahan, J.E., Short, J.M., Short, H., Bruzel, A., Wynshaw-Boris, A., Meisner, H.M., Samols, D., and Hanson, R.W. (1983) Isolation and Characterization of the Gene Coding for Cytosolic Phosphoenolpyruvate Carboxykinase (GTP) from the Rat. *Proc. Natl. Acad. Sci. U.S.A.*, 80:3656-3660.
2. Wynshaw-Boris, A., Lugo, T.G., Short, J.M., Fournier, R.E.K., and Hanson, R.W. (1984) Identification of cAMP Regulatory Region in the Gene for Rat Cytosolic Phosphoenolpyruvate Carboxykinase (GTP): Use of Chimeric Genes Transfected into Hepatoma Cells. *J. Biol. Chem.*, 259:12161-12169.
3. Wynshaw-Boris, A., Lugo, T.G., Short, J.M., Fournier, R.E.K., and Hanson, R.W. (1985) A Region of the Gene for Rat Cytosolic P-enolpyruvate Carboxykinase Confers cAMP Responsiveness to the HSV-thymidine Kinase Gene. In: *Membrane Receptors and Cellular Recognition*, (M. Czech and C.R. Kahn, eds.), Alan Liss Inc., New York, pp 339-346.
4. Wynshaw-Boris, A., Short, J.M., and Hanson, R.W. (1986) Characterization of the Phosphoenolpyruvate Carboxykinase (GTP) Promoter-Regulatory Region. I. Multiple Hormone Regulatory Elements and the Effects of Enhancers. *J. Biol. Chem.*, 261:9714-9720.
5. Short, J.M., Wynshaw-Boris, A., Short, H.P., and Hanson, R. W. (1986) Characterization of the Phosphoenolpyruvate Carboxykinase (GTP) Promoter-Regulatory Region. II. Identification of cAMP and Glucocorticoid Regulatory Domains. *J. Biol. Chem.*, 261:9721-9726.
6. Wynshaw-Boris, A., Short, J.M., and Hanson, R.W. (1986) The Determination of Sequence Requirements for Hormonal Regulation of Gene Expression. *Biotechniques*, 4:104-119.
7. Burns, D.M., Bhandari, G., Short, J.M., Sanders, P.G., Wilson, R.H., and Miller, R.E. (1986) Selection of a Rat Glutamine Synthetase cDNA Clone. *Biochemical and Biophysical Research Communications*, 134:146-151.
8. Hod., Y. Cook, J.S., Weldon, S.L., Short, J.M., Wynshaw-Boris, A., and Hanson, R.W. (1986) Differential Expression of the Genes for the Mitochondrial and Cytosolic Forms of P-enolpyruvate Carboxykinase Gene. In: *Metabolic Regulation: Application of Recombinant DNA Techniques*, (A.G., Goodridge and R.W. Hanson eds.), Annals of the New York Academy of Sciences, New York, Vol. 278, pp. 31-45.
9. Wynshaw-Boris, A., Short, J.M., and Hanson, R.W. (1987) *cis* - acting Regulatory Elements in Horizontally Responsive Genes. In: *Progress in Nucleic Acid Research and Molecular Biology* (W.E. Cohn and K. Moldave eds.), Academic Press, Inc., Orlando, Florida, 34:59-87.

10. Bullock, W., Fernandez, J.M., and Short, J.M. (1987) XL1-Blue: A High Efficiency Plasmid Transforming *recA E.coli* Strain With β -Galactosidase Selection. *Biotechniques*, 5:60-64.
11. Short, J.M., Fernandez, J.F., Sorge, J.A., and Huse, W. (1988) Lambda ZAP[®]: A Bacteriophage Lambda Expression Vector With *In Vivo* Excision Properties. *Nucleic Acids Res.*, 16:7583-7600.
12. Short, J.M. (1988) Book Review: Vectors - A Survey of Molecular Cloning Vectors and Their Uses. Raymond L. Rodriques and David T. Denhardt, eds, Butterworths, Stoneham, MA. *Genomics*, 2:270-271.
13. Short, J.M., and Pollard, A. (1988) Gigapack XL: Size Selective Packaging Extract. *Strategies in Mol. Biol.*, 1:5-7.
14. Kretz, P.L., and Short, J.M. (1989) Gigapack II: A Restriction Deficient (*mcrA*-, *B*-, *hsd*-, *mrr*-) Lambda Packaging Extract. *Strategies in Mol. Biol.*, 2(2):25-26.
15. Kretz, P.L., Reid, C.H., Greener, A., and Short, J.M. (1989) Effect of Lambda Packaging Extract M_{cr} Restriction Activity on DNA Cloning. *Nucleic Acids Res.* 17:5409.
16. Sastry, L., Alting-Mees, M., Huse, W.D., Short, J.M., Sorge, J.A., Hay, B.N., Janda, K.D., Benkovic, S.J., and Lerner, R.A. (1989) Cloning of the Immunological Repertoire in *E. coli* for Generation of Monoclonal Catalytic Antibodies I. Construction of a V_H Specific cDNA Library. *Proc. Natl. Acad. Sci. U.S.A.*, 86:5728-5732.
17. Short, J.M. (1989) The Use of Lambda Phage Shuttle Vectors in Transgenic Mice for Development of a Short Term Mutagenicity Assay. In: *Fifth International Conference on Environmental Mutagens*, Alan Liss, Inc., New York, Part A:335-367. Article and Lecture.
18. Alting-Mees, M., and Short, J.M. (1989) pBluescript II: Gene Mapping Vectors. *Nucleic Acids Res.*, 17:9494.
19. Shopes, B., Alting-Mees, M., Amberg, J.R., Ardourel, D., Callahan, M., Detrick, J., Hay, B.N., Hogrefe, H.H., Greener, A., Gross, E.A., Kubitz, M.M., Mullinax, R.L., Wilson, C., Short, J.M., and Sorge, J.A. (1990) Bacteriophage Immuno-expression Libraries: An Emerging Technology for the Identification and Production of Monoclonal Antibodies. *Antibody Engineering, New Tech. & Application Implications*. pp. 98-101.
20. Alting-Mees, M., Amberg, J., Ardourel, D., Elgin, E., Greener, A., Gross, E.A., Kubitz, M., Mullinax, R.L., Short, J.M., and Sorge, J.A. (1990) Monoclonal Antibody Expression Libraries: A Rapid Alternative to Hybridomas. *Strategies in Mol. Biol.*, 3:1-9.
21. Kohler, S., Provost, S., Dyaico, M., Sorge, J., and Short, J.M. (1990) Development of a Short-term, *In Vivo* Mutagenesis Assay: The Effects of Methylation on the Recovery of a Lambda Phage Shuttle Vector from Transgenic Mice. *Nucleic Acids Res.*, 18:3007-3013.
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